

Transmitted herewith is a certified copy of foreign application 2,218,199 filed December 9, 1997 in Canada.

In the Claims

Please amend Claims 1, 3-5, 7-9 and 11-12 as follows. Amendments to the claims are indicated in the attached "Marked Up Version of Amendments" (pages i – ii).

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- C 1. (Twice Amended) An isolated human PAB II gene comprising a polymorphic GCG repeat in exon I thereof, wherein said polymorphic GCG repeat has the sequence ATG (GCG)<sub>6+n</sub> GCA,  
with n being selected from 1 to 7 and wherein said polymorphic repeat of said GCG repeat is indicative of a disease in a human patient.

- C2 3. (Twice Amended) The gene of claim 1, wherein n is selected from 2 to 7, and wherein said polymorphic repeat of said GCG repeat is associated with an increased severity of the disease.

4. (Amended) The gene of claim 3, wherein a phenotype associated with said polymorphic repeat of said GCG repeat is dominant.

5. (Twice amended) The gene of claim 1, wherein a first allele of said GCG repeat of said human patient has an n which is equal to 1.

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- C3 7. (Amended) The gene of claim 1, wherein said gene is isolated from a patient who is homozygous for said polymorphic GCG repeat.

8. (Amended) The gene of claim 1, wherein said gene is isolated from a patient who is heterozygous for said polymorphic GCG repeat.